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Disturbed wnt signalling due to a mutation in CCDC88C causes an autosomal recessive non-syndromic hydrocephalus with medial diverticulum

Ekici, A B ; Hilfinger, D ; Jatzwauk, M ; Thiel, C T ; Wenzel, D ; Lorenz, I ; Boltshauser, E ; Goecke, T W ; Staatz, G ; Morris-Rosendahl, D J ; Sticht, H ; Hehr, U ; Reis, A ; Rauch, A

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Disturbed Wnt Signalling due to a Mutation in *CCDC88C* Causes an Autosomal Recessive Non-Syndromic Hydrocephalus with Medial Diverticulum

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Key Words

Brain malformation · *CCDC88C* · Daple · Hydrocephalus · Wnt signalling

Abstract

The etiology of non-syndromic hydrocephalus is poorly understood. Via positional cloning in a consanguineous family with autosomal recessive hydrocephalus we have now identified a homozygous splice site mutation in the *CCDC88C* gene as a novel cause of a complex hydrocephalic brain malformation. The only living patient showed normal psychomotor development at the age of 3 years and 3 months and her deceased aunt, who was assumed to suffer from the same condition, had mild mental retardation. The mutation in the affected patients, a homozygous substitution in the donor splice site of intron 29, resulted in a shorter transcript due to exclusion of exon 29 and loss of functional protein, as shown by Western blotting (p.S1591HfsX7). In normal human tissue panels, we found *CCDC88C* ubiquitously expressed,

but most prominently in the fetal brain, especially in pons and cerebellum, while expression in the adult brain appeared to be restricted to cortex and medulla oblongata. *CCDC88C* encodes DAPLE (HkRP2), a Hook-related protein with a binding domain for the central Wnt signalling pathway protein Dishevelled. Targeted quantitative RT-PCR and expression profiling of 84 genes from the Wnt signalling pathway in peripheral blood from the index patient and her healthy mother revealed increased mRNA levels of *CCDC88C* indicating transcriptional upregulation. Due to loss of *CCDC88C* function β -catenin (*CTNNB1*) and the downstream target *LEF1* showed increased mRNA levels in the patient, but many genes from the Wnt pathway and transcriptional target genes showed reduced expression, which might be explained by a complex negative feedback loop. We have thus identified a further essential component of the Wnt signalling pathway in human brain development.

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A prenatal hydrocephalus is observed in about 1:2,000 pregnancies. The etiology is heterogeneous including intrauterine infections, neural tube defects, chromosomal aberrations and a wide variety of monogenic syndromes, twinning as well as CNS tumors, but remains unsolved in the majority of non-syndromic cases [Garne et al., 2010]. Prognosis critically depends on the presence of additional cerebral and extracranial malformations, which are observed in about 70–80% of the cases [Schrander-Stumpel and Fryns, 1998]. Important syndromal forms of congenital hydrocephalus include the autosomal recessive syndromal lissencephalies type 2 Walker-Warburg syndrome and Muscle-eye-brain disease with additional observation of a hypoplastic cerebellum and brainstem together with cobblestone lissencephaly and agenesis or hypoplasia of the corpus callosum as well as postnatal congenital muscular hypotonia, various eye abnormalities and global developmental delay, which in about half of the cases result from mutations in one of currently 6 genes associated with the O-glycosylation of α -dystroglycan [Bouchet et al., 2007]. The most frequent monogenic form of congenital hydrocephalus is due to hemizygous mutations in the *LICAM* gene (about 7–15%) and clinically characterised by the distinct combination of a hydrocephalus due to stenosis of the aqueduct of Sylvius commonly in association with adducted thumbs (HSAS) [Tapanes-Castillo et al., 2010; Vos et al., 2010].

For siblings of sporadic patients with non-syndromic hydrocephalus a recurrence risk of about 1–4% has been determined and together with several monogenic animal models reflects the contribution of genetic factors to the pathogenesis of hydrocephalus [Schrander-Stumpel and Fryns, 1998; Zhanget al., 2006]. However, besides *LICAM*, no further genes underlying human non-syndromic congenital hydrocephalus have yet been identified. By autozygosity mapping, we now revealed a homozygous mutation in the *CCDC88C* gene as underlying an autosomal recessive inherited form of non-syndromic complex hydrocephalus with apparently normal or at most mildly retarded psychomotor development.

Patients and Methods

Patients

The study was approved by the ethical review board of the medical faculty of the Friedrich-Alexander University of Erlangen-Nuremberg and informed consent was obtained from the parents. We studied a consanguineous family; 58 patients with schizencephaly, 20 of which had additional brain malformations including 3 patients with solitary cerebral cysts, and 5 patients

with variable brain malformations with enlarged cerebrospinal fluid spaces.

Family History

The simplified pedigree of this consanguineous family is depicted in figure 1A. IV:1, a healthy woman of Algerian origin presented during her first pregnancy at 26 weeks of gestation with abnormal fetal ultrasound showing enlarged cerebral ventricles (fig. 1B). Fetal karyotyping from amniotic fluids revealed a normal female karyotype at a 500 GTG-banding resolution. At 29 weeks of gestation schizencephaly was suspected and after exclusion of maternal contamination using 13 microsatellite markers (PowerPlex16; Promega), *EMX2* sequencing was performed on fetal DNA, which showed normal results. The pregnancy was eventually terminated and no post-mortem examination was performed. Similar ultrasound findings were detected in the second pregnancy of IV:1 at 25 weeks of gestation, but this time the parents decided to give birth to the girl. V:2 was born at 38 weeks of gestation with a weight of 2,490 g, a length of 45 cm and a head circumference of 34 cm. Apgar scores were 9-10-10, umbilical cord pH was 7.18 and base excess was –8. Cerebral MRI scans at the age of 3 days revealed marked dilatation of the lateral ventricles, while the third and fourth ventricles were not enlarged (fig. 1C, D, E). No cerebrospinal fluid flow void was evident in the aqueduct which may indicate either primary or secondary stenosis. Since no follow-up MRI was available, this cannot be further evaluated. A diverticulum-like pouch extended from the medial-posterior aspect of the left lateral ventricle into the interhemispheric space, extending through the tentorium into the infratentorial space, leading to mild compression of the upper cerebellar vermis. The posterior fossa was markedly enlarged with supra- and retrocerebellar fluid accumulation. The vermis was of normal shape, not elevated, and the communication between the fourth ventricle and the cisterna magna was of normal dimension. Brainstem morphology was not remarkable. Electroencephalography at the age of 5 days showed bitemporal and mild central hypersynchronous activity with sharp waves and spikes. The child showed no other malformations, dysmorphism or neurological anomalies. Because of increasing head circumference (40.3 cm) a ventriculo-peritoneal shunt was placed at the age of 4 weeks in the absence of any signs of increased intracranial pressure. At the age of one year the girl was hospitalized because of a tonic-clonic seizure with fever of 40.2°C after a few days of gastroenteritis with diarrhea and vomiting. At the age of 12.5 months she was found normally developed. She was able to walk assisted and produced sounds. Body weight was 9.4 kg (50th centile), height was 72.5 cm (10th centile) and head circumference was 45.5 cm (25–50th centile). At the age of 3 years and 3 months she spoke Arabic and German in 3-word sentences with grammatical errors and dyslalia. Communication, playing behaviour, coordination and fine motor skills were normal. Bayley Scales of Infant development testing revealed 156 points corresponding to a developmental quotient of 95. She had convergent strabismus which was treated with plasters, which she did not tolerate well. Her height was 94 cm (10th centile), her weight was 13 kg (10th centile) and her head circumference was 48.7 cm (25th–50th centile).

The family history was remarkable for parental consanguinity (first cousins once removed). The mother of V:I and V:II, who's parents were first cousins, in addition to 4 healthy siblings had two sisters who died neonatally of unknown reason. A further

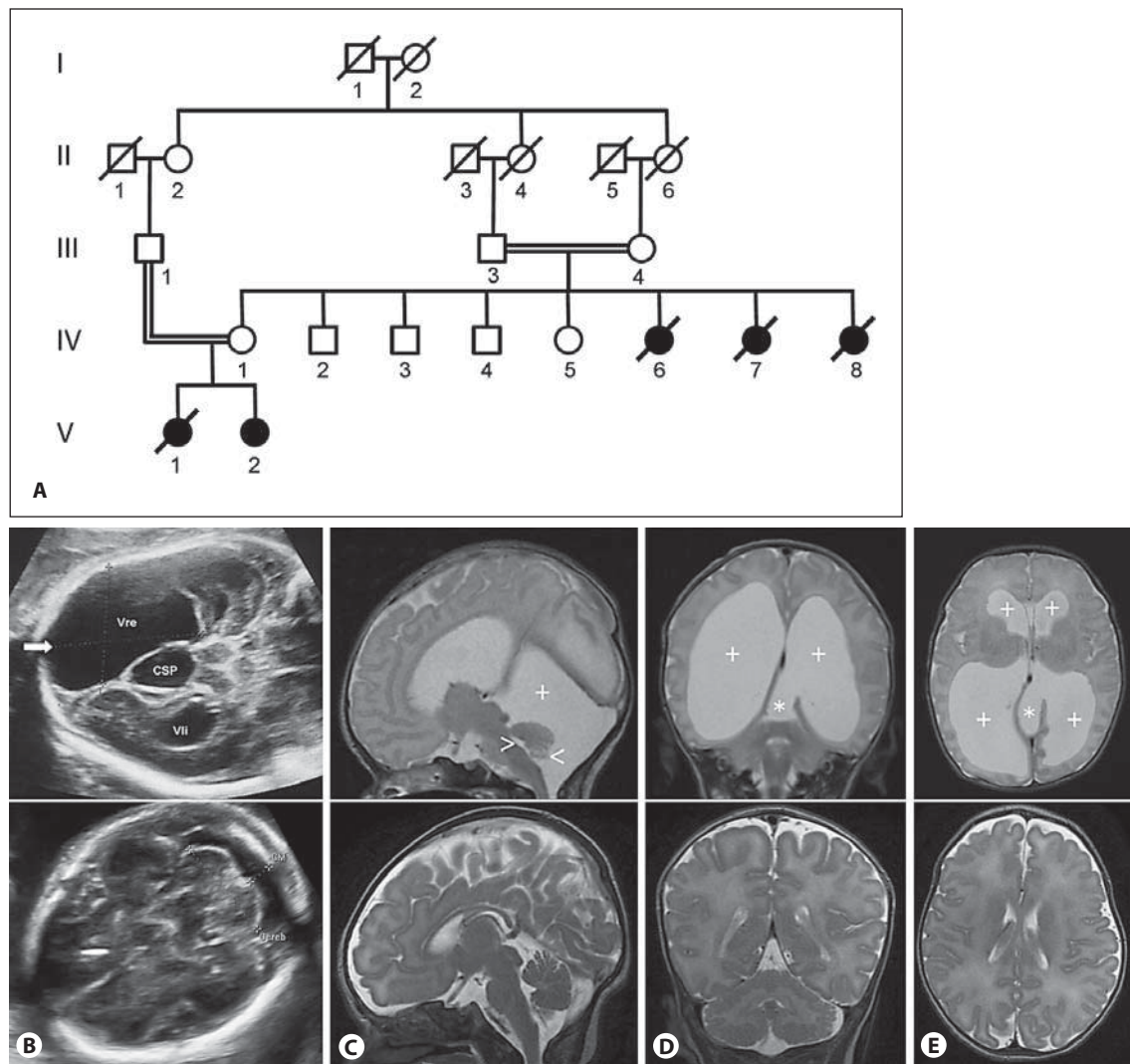


Fig. 1. Pedigree and clinical findings in the investigated family. **A** Pedigree of the consanguineous family. IV:6 and IV:7 died neonatally of unknown cause. IV:8 died at the age of 22 years and was reported to have suffered from mild mental retardation. V:1 and V:2 showed a similar hydrocephalic brain malformation. III:1, IV:1, V:1 and V:2 were clinically investigated by the authors. **B–E** Ultrasound and MRI in affected individuals and normal scans for comparison in the lower row. **B** Ultrasound examination of V:1 at 29 weeks of gestation, demonstrating marked ventricular dilatation and an interhemispheric cystic structure in the posterior aspect (CSP). **C, D, E** T2w MRI of V:2 at 5 days. **C** Midsagittal

MRI demonstrating a small vermis in normal position (<), a small 4th ventricle (>), marked retrocerebellar fluid accumulation in an enlarged posterior fossa (+), and no flow void in the aqueduct (data not shown). **D** Coronal section demonstrating pronounced ventricular dilatation (+). There is a diverticular-like extension from the medial aspect of the left lateral ventricle (trigone) extending through the tentorium into the posterior fossa (*). **E** Axial view demonstrating marked dilatation of the lateral ventricles (+). The diverticular-like extension is visible at the medial aspect of the left trigone (*). The lower row of ultrasound and MRI figures represents comparable normal sections.

sister IV:8 died at the age of 22 years. She was reported mentally retarded with a developmental age of 8–10 years.

Patient Samples

Genomic DNA from whole EDTA blood, saliva or amniotic fluid samples was extracted using a Flexigene AGF3000 technology (Qiagen, Hilden, Germany) on an automated Autogenflex

3000 isolator (Autogen, Holliston, Mass., USA) according to the protocols supplied by the provider. cDNA was synthesized from RNA extracted from fresh blood samples collected in PAXgene tubes (Becton Dickinson, Heidelberg, Germany) using the Superscript II Reverse Transcriptase Kit with random hexamer primers (Invitrogen, Carlsbad, Calif., USA).

Table 1. Primers used for sequencing *CCDC88C*

Primer forward	Sequence	Primer reverse	Sequence
5'UTR_p1_f	agcgaatggcacacactcc	5'UTR_p1_r	gtacaggggtgacccgaagg
5'UTR_p2_f	agcgcctgggctgtaatacc	5'UTR_p2_r	caggaagcaggtgatgtgg
5'UTR_p3_f	ttgcccttttctgtcg	5'UTR_p3_r	ctccgaggtgctctgc
5'UTR_p4_f	ggggcctcttgactgtacc	5'UTR_p4_r	ctcggggaggagaacaatacg
Ex1n_f	cctcgggccaccagtc	Ex1n_r	ggctccagagtacaaagc
Ex2_f	tcctcctgtctcgtgtctgtt	Ex2_r	cgtctccaccttcagagaactac
Ex3_f	ctgcactgatattgtgttggtt	Ex3d_r	cctaatacataagcaccgtttcc
Ex3b_p1_f	caccctgtaaggagaaggg	Ex3b_p1_r	cctccaggggatcacacg
Ex3b_p2n_f	atcactgtgggctcttctgg	Ex3b_p2n_r	tggtactggtcccttgacagg
Ex3b_p3_f	actgcaacctcatcctctgc	Ex3b_p3_r	atcccttctccactgatgg
Ex3b_p4n_f	gcaagttttgctgttctgc	Ex3b_p4n_r	gctagggccttgacaataacc
Ex4_f	gtttgcatcttacgtgctcctt	Ex4_r	ctcagggtaatagaaccactctt
Ex5_f	tggaagatcgatggagtgtgta	Ex5_r	gtactagaaacagctgccacgtaa
Ex6_f	agacacatcagtgtaaggctgg	Ex6_r	caatgtgtctgtgtgtctgaa
Ex7_f	ctgaaagcagaagtgttctgta	Ex7_r	ctgaagatgaaggagaggagat
Ex8_f	accttctcctcgtttcttagtc	Ex8_r	agctagtcagggtcaagagta
Ex9_f	agtggcggatggacatagtta	Ex9_r	atctccacctgctgtcactaaga
		Ex10_i_r	aatttcacagacccccatc
		Ex10_a_r	ggagaaacccctgaatcactgc
Ex10_f	cctgtttcttagtgacagcaggt	Ex10_r	acaggctcagtctgaatgtgctt
Ex11_f	ctcggacttcctctagattgtt	Ex11_r	caagtagatggaactacaggctca
Ex12_f	agtggagcagtagatcttcatcc	Ex12_r	caagtgagccttgaaaggaaatc
Ex13_f	tgatgacctctcggagctt	Ex13d_r	tgacacctgcatgctctacc
Ex13bf	gtgggttctggcataagacc	Ex13br	tgtagcaactccccacacc
Ex14_f	gctacagaatgatgtcaggaggt	Ex14_r	cagtaccaggaatggctgtaatc
Ex15_p1_f	tctaagtgggtgtgtctgtgc	Ex15_p1_r	ttctcgggtgctgtctcc
Ex15_p2_f	agctgcacaggagacttg	Ex15_p2_f	ctcgtctccaggctctgc
Ex15_p3_f	gaggaagaacgtggatcagcgctcaagg	Ex15_p3_r	gggtgactgctgtgtgagg
Ex15_p4_f	aggtggagctcaaggatgc	Ex15_p4d_r	aaagcatttaaggaggacacaga
Ex16_f	atgtagccacctggtctggtt	Ex16_r	actgaacagtgagggaagtgtct
Ex17_f	gaacttgcggagaggaagtagat	Ex17_r	gtatcttgcctgtgtgacttg
Ex18_f	cgtctggcttacacagacca	Ex18d_r	aggacctgcaactcaagacg
Ex19_f	agtacctgtggccagtagcat	Ex19_r	gatcactgtttacagcgggtct
Ex20_f	gtaccctctgggtcttctgc	Ex20_r	caggccccacctttcc
Ex21_f	ctctgggatgtgtgactgtgc	Ex21_r	ggacttgacttctcagttgttg
Ex22_f	gttacctgtttgtcaggctcttc	Ex22_r	cctccacacatttctgtctttg
Ex23_f	ttcttcttgggagctaggtt	Ex23_r	agaagtttctgtgtgtgaaag
Ex24_f	ttccaaactcagtgacctagac	Ex24d_r	gattctgaactctctgtatgtgg
Ex25_f	aacatcccagcccatgttag	Ex25_r	ctcgtaacacatctaagcgaagg
Ex25b_f	caccagctctctctaaacg	Ex25b_r	ttttaggaagtgcagcaacc
Ex26b_f	cctctgacaccacctgaagc	Ex26b_r	agactgagaggagagcaacc
Ex27_f	agcagaggtggaagtgtgattt	Ex27_r	cttaaggaaaggtaccgtgga
Ex27b_f	tccgtaacctgactcactagc	Ex27b_r	tcgaatccaatgacaagtgc
Ex28_f	ctgggatatacctggtcattgg	Ex28_r	gtagcatgatcacagcccactat
Ex29_f	gaagcctctagaagccacca	Ex29n_r	ttgcgggtgactgtcagcta
Ex30_p1n_f	acatgtgcctcggagaagg	Ex30_p1n_r	ctgaggggactcctgtttgc
Ex30_p2_f	aagaaggggccaagatgc	Ex30_p2d_r	ggagctgccagcctttcc
Ex30_p3_f	cctgtctctgggcagacc	Ex30_p3_r	ctgcagcaggtgagaagtgc
Ex30_p4_f	gacacgaggcgcttctcc	Ex30_p4d_r	cggactgctctttgagacg
Ex30_p5_f	gtcggctcctccgatttgg	Ex30_p5_r	ttgcaaatgagtgttctactgg
3'UTR_p1_f	tgaacgtgtttctgtgtctgc	3'UTR_p1_r	ctgtgtgtgtcccaaac
3'UTR_p2_f	aagtgtgaggtgctgtgttcc	3'UTR_p2_r	gaatcttgaccaaagacacaaagc
3'UTR_p3_f	gggtcatccgcacagagc	3'UTR_p3_r	acccgatgctaacaatgtgc
3'UTR_p4_f	ctgctgcttgtgtcttgg	3'UTR_p4_r	gcaggagagcccaagtc

Table 2. Primers used for isoform-specific RT-PCR of *CCDC88C*

Primer forward	Sequence	Primer reverse	Sequence
RT_IF1f	aaagccaggcagatcaagg	RT_IF1r	aggcgctctgacttcttcg
RT_IF2f	gccaggatgtccgtactaagc	RT_IF2r	cgaggcaaggggtactcg
RT_IF3af	cagcctcatttctcaatgtcc	RT_IF3ar	tgcttgaaggcatgaagagg
RT_IF3bf	ctgctgccctgcttctagg	RT_IF3br	tgctggatgtgttctactgc
RT_IF3f	ctacagccctccaactcc	RT_IF3r	ctgggatcaggtcttcg
RT_IF4nf	ggacggcatcttttgaacc	RT_IF4nr	aatgcagacactcccagacc

Linkage Analysis

We performed genome-wide linkage analysis and homozygosity mapping in the consanguineous family (fig. 1A). All living individuals shown in the pedigree were available for linkage analysis. In addition DNA from amniotic fluid of V:1 was included in the genetic studies. Thus, DNA samples of the 2 affected children, both unaffected parents and seven further unaffected relatives were genotyped with the GeneChip Human Mapping 250K Nsp Array (Affymetrix, Santa Clara, Calif., USA). Genotypes were called by the GeneChip Genotyping software (GTTYPE v4.0; Affymetrix). Simulation analysis for expected LOD scores (ELOD) for the given family pedigree was performed with a parametric 2-point analysis with the software FastSLink [Weeks et al., 1990]. Relationships of family members were verified by checking the genotype data with the software Graphical Representation of Relationships (GRR) [Abecasis et al., 2001]. Mendelian errors due to genotype errors were checked with the software Pedcheck [O'Connell and Weeks, 1998] and erroneous genotypes were removed from the data file. Parametric multipoint linkage analysis and haplotype construction were done with the programs Allegro [Gudbjartsson et al., 2000, 2005] and Merlin [Rampersaud et al., 2005]. Data handling was done with the software easyLinkage-Plus [Hoffmann and Lindner, 2005] under the assumption of an autosomal recessive mode of inheritance with 100% penetrance, disease allele frequency of 0.01% and at least 0.01 cM intermarker distance. Visualisation of haplotype data was performed using the software haploPainter [Thiele and Nürnberg, 2005].

Mutation Analysis of *CCDC88C*

Primers for all 30 exons, the complete 3' and 5' UTR, including 1,500 bp flanking promotor region were designed with the software ExonLocator [Schageman et al., 2004] and Primer3 [Rozen and Skaletsky, 2000]. Primer sequences are summarized in table 1. Genomic DNA of participants was whole genome amplified with Phi29 polymerase (GenomiPhi v2; GE Healthcare, UK) for mutation analysis. Subsequently all sequence aberrations were verified with original genomic DNA. Genomic DNA (20–50 ng) was amplified by PCR in 20-μl reactions with intronic primers according to standard procedures. The PCR products were purified with the AMPure Kit (Agencourt, Beverly, Mass., USA) on a Biomek NX instrument (Beckman Coulter, Fullerton, Calif., USA). Sequencing reactions were performed on both strands using the BigDye Terminator Cycle Sequencing Kit v3.1 (Applied Biosystems, Foster City, Calif., USA) according to the manufacturer's instructions. After purification using the CleanSEQ Kit (Agencourt), the products were analyzed on an ABI Genetic Ana-

Table 3. Exon localisation and expected fragment length of isoform-specific primers of *CCDC88C*

Product	Starting exon	Ending exon	Length, bp
IF1	14/15	15	617
IF2	25b	29	488
IF3a	26b	27b	403
IF3f3ar	26	27b	234
IF3b	26b	28	540
IF4	2	3b	540

lyzer 3730 (Applied Biosystems) and the traces were evaluated using the SeqPilot 3.2.1.5 software (JSI medical systems, Costa Mesa, Calif., USA).

Characterization of Splicing Pattern of Intron 29

To ascertain the effect of the IVS29 + 1 splice site mutation we amplified exons 27–30 and 28–30 from cDNA to generate wt amplicons of 435 bp and 541 bp, respectively. Amplified PCR products were evaluated by sizing in agarose gels and direct sequencing.

Western Blotting

Western blot analysis using protein lysates extracted from Epstein-Barr virus-transformed lymphoblastoid cell lines (LCLs) of patient V:II, her mother and controls were performed using a rabbit monoclonal KIAA1509 (*CCDC88C*) antibody which detects aa1566 to 1693 (encoded by exons 28 and 29) (Sigma Aldrich, St. Louis, Mo., USA). A β-actin antibody (ab8227) was used as control and a HRP-conjugated goat anti-rabbit IgG antibody (170–6515; Bio-Rad, Munich, Germany) as secondary antibody. Electrophoresis was performed on a 3–8% Tris-Acetate gel using the NuPage Novex system from Invitrogen.

Expression Studies of *CCDC88C* in Human Tissue Panels

The expression profile of *CCDC88C* was investigated by isoform specific RT-PCR (tables 2 and 3) and quantitative RT-PCR with the *CCDC88C* pre-designed TaqMan Gene Expression Assay Hs00929473_m1 (Applied Biosystems) using following cDNA panels: Human Multiple Tissue cDNA (MTC) Panel I (Clontech), Human Fetal Multiple Tissue cDNA (MTC) Panel (Clontech), Human Adult Neural Tissue II (BioChain) and Human Fetal Neural Tissue II (BioChain).

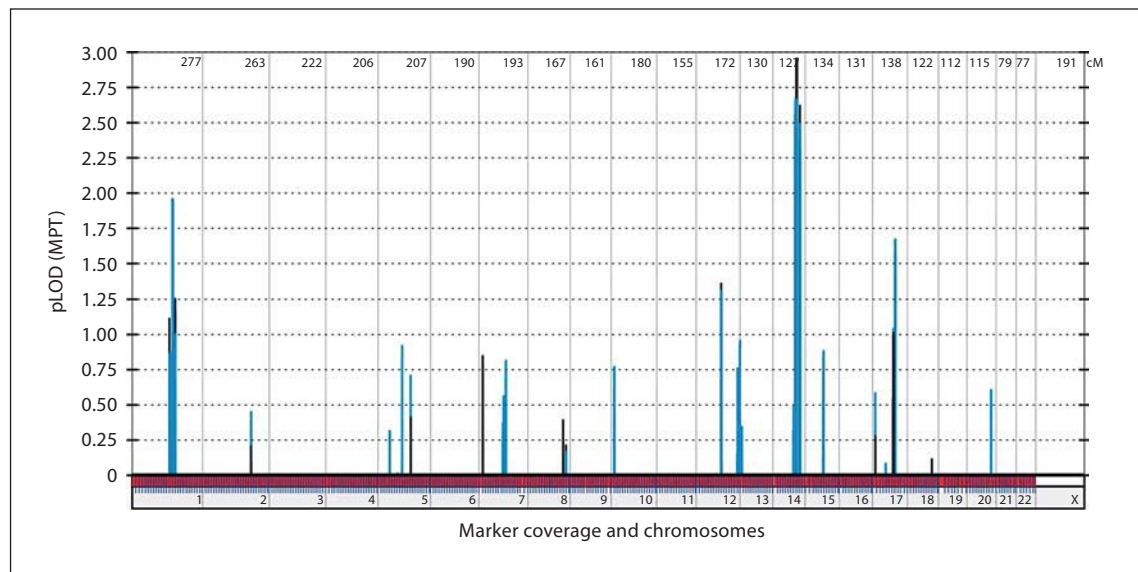


Fig. 2. Results of whole genome linkage analysis. Parametric multipoint LOD-scores of pedigree subset A are in blue and those of subset B are in black. Both pedigree parts indicate linkage to a single locus on chromosome 14 with a maximum LOD-score close to 3. As both subsets coincide but could not be calculated together we assume that the overall LOD-score clearly reaches genome-wide significance.

Expression Studies in Patient Samples

To measure the expression of *CCDC88C* and β -catenin pre-designed TaqMan Gene Expression Assays were used (Hs00929473_m1, Hs00170025_m1; Applied Biosystems). Expression levels in patients' RNA were quantified by real-time PCR in 4 replicates each as previously described [Rauch et al., 2008] and results were normalized to the mean of 3 endogenous controls (β -2-microglobulin (huB2M), β -actin (huACTB), acidic ribosomal protein (huPO)). Normalized expression levels were set in relation to 8 controls.

Expression levels of 84 genes from the Wnt pathway were analysed using the Wnt Signaling Pathway PCR Arrays (SABioscience, Frederick, Md., USA) which is based on quantitative RT-PCR using SYBR green detection on an ABI PRISM 7900HT device (Applied Biosystems) according to the supplier's instructions. Six different healthy, unrelated controls were analysed once, while the patient and her mother were analysed four times each.

Results

Linkage and Mutation Analysis

Whole genome LOD score analysis under the assumption of full penetrance and a disease allele frequency of 0.01% revealed a maximum LOD score of 2.96 on chromosome 14q (fig. 2). This finding was in line with the simulation analysis and the obtained ELOD score of 2.56. Due to the complexity of the pedigree, though, it was impossible to calculate LOD-scores using the computer program AL-

legro on any of our computers. Only when we portioned the pedigree we were able to perform the analysis. For the first calculation we excluded the healthy individuals IV:2 and IV:3, for the second IV:4 and IV:5. Both calculations indicated linkage to a single locus on chromosome 14 with a maximum LOD-score close to 3. As both subsets coincided but could not be calculated together we assume that the overall LOD-score clearly reaches genome-wide significance. Haplotype analysis revealed a linked region of 3.37 Mb on chromosome 14 flanked by the markers SNP_A-1934157 (rs41463644) and SNP_A-1840441 (rs7148382) (fig. 3) containing 25 positional candidate genes, including all RefSeq, orthologous and predicted genes. In the analysis for potential candidate genes, we focused on genes which were expressed in the brain and/or found in signal transduction pathways in which genes are involved in cerebral cortex malformations [Brunelli et al., 1996; Granata et al., 1997; Cecchi, 2002; Theil et al., 2002; Muzio et al., 2005]. The most promising gene was *CCDC88C* (NM_001080414). The gene encompasses 30 exons and codes for the DVL-binding protein DAPLE. It interacts as a homooligomer with the PDZ domain of DVL1 and functions as a negative regulator of the canonical Wnt signaling pathway to inhibit CTNNB1/ β -catenin stabilization [Oshita et al., 2003]. By direct sequencing of *CCDC88C* exons of the affected child V:2 we identified a homozygous

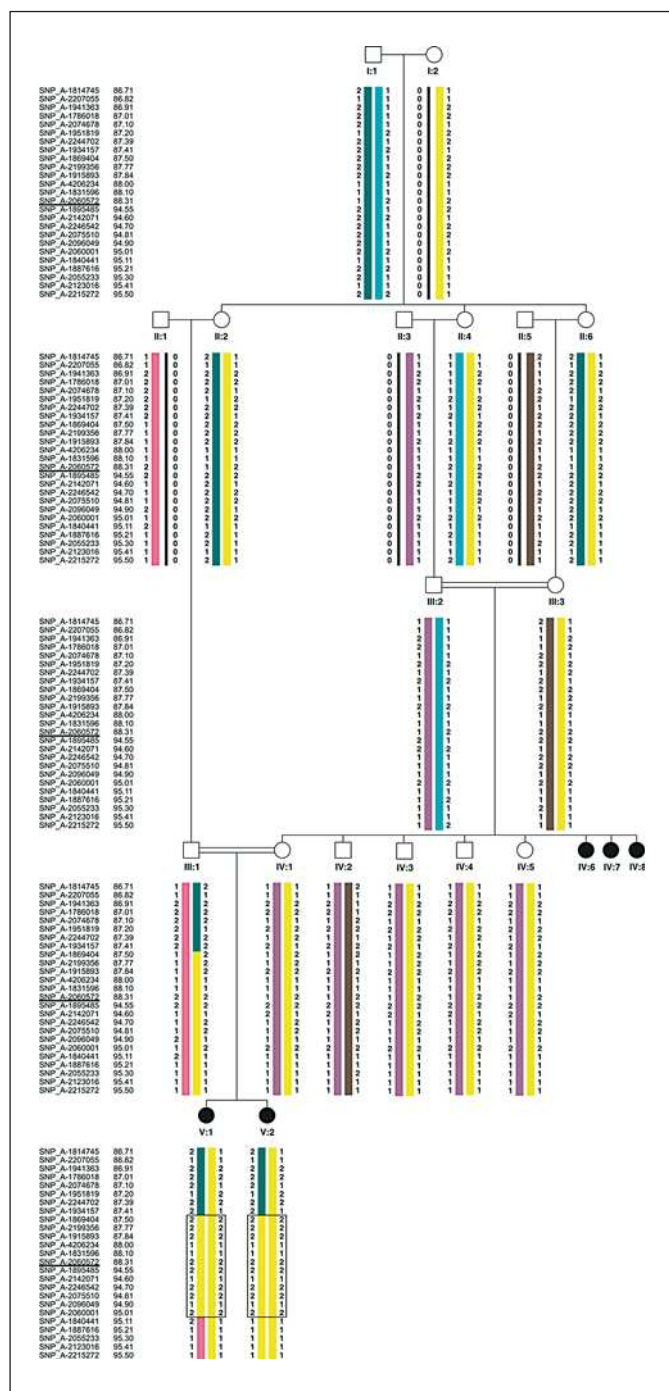


Fig. 3. Haplotypes of the linked interval in the pedigree. The linked region of 3.37 Mb on chromosome 14 is flanked by the markers SNP_A-1934157 (rs41463644) and SNP_A-1840441 (rs7148382).

splice site mutation at position IVS29 + 1 which changes the splice site donor G-U to A-U (c.5058 + 1G>A). The affected fetus V:1 was also homozygous for this mutation, while, as expected, both parents were heterozygous. All other unaffected family members were homozygous for the wild-type allele or heterozygous (fig. 4A). The mutation was not detected in 224 German healthy controls.

To further evaluate the splice site mutation we amplified two different exon 29 spanning fragments (exons 27–30 and 28–30) from the cDNA of the affected child V:2 and the mother IV:1. The mother and a control sample showed solely the wt amplicons of 435 bp and 541 bp, respectively, by sizing in agarose gel. Shortened amplicons were detectable only in the affected child V:2 (fig. 4B). Lack of amplification of the abnormal transcript in the mother may be explained by privileged amplification of the wild-type allele which is absent in the patient. By direct sequencing of the shortened transcript we could confirm the deletion of 290 bp corresponding to exon 29. Since exon 29 does not end with a codon this results in a frame shift in exon 30 and destroys the Dishevelled binding site which is encoded by the 3 terminal codons (p.S1591HfsX7). Western blotting confirmed the absence of wild-type protein in the patient, but could not exclude the presence of an abnormal protein, since the antibody binds to the amino acids encoded by the deleted exon 29 (fig. 4C).

While none of the 58 patients with schizencephaly showed a potential mutation, one of the 5 patients with enlarged cerebrospinal fluid spaces, abnormal gyral pattern and profound developmental delay was found to harbour two compound heterozygous variants. The transition c.1715C>T in exon 15 leads to the amino acid exchange p.S572L in a highly conserved position. The transversion c.5954C>G (rs3814839) in exon 30 also results in an amino acid exchange (p.S1985C). While the exon 30 variant was not detected in 504 controls, the variant in exon 15 was detected twice in heterozygous status. The parents were heterozygous for one of both variants, respectively, but the healthy sister was heterozygous for both mutations. Therefore the mutations were not considered to be the cause of the disease in the patient.

Expression Studies in the Patient and Her Heterozygous Mother

Quantitative RT-PCR studies in RNA derived from PAXgene samples of patient V:2 and her heterozygous carrier mother revealed an 8.3-fold increased *CCDC88C* expression in the patient and an 3.7-fold increased expression in the mother. These findings confirm the absence of nonsense mediated mRNA decay and indicate increased

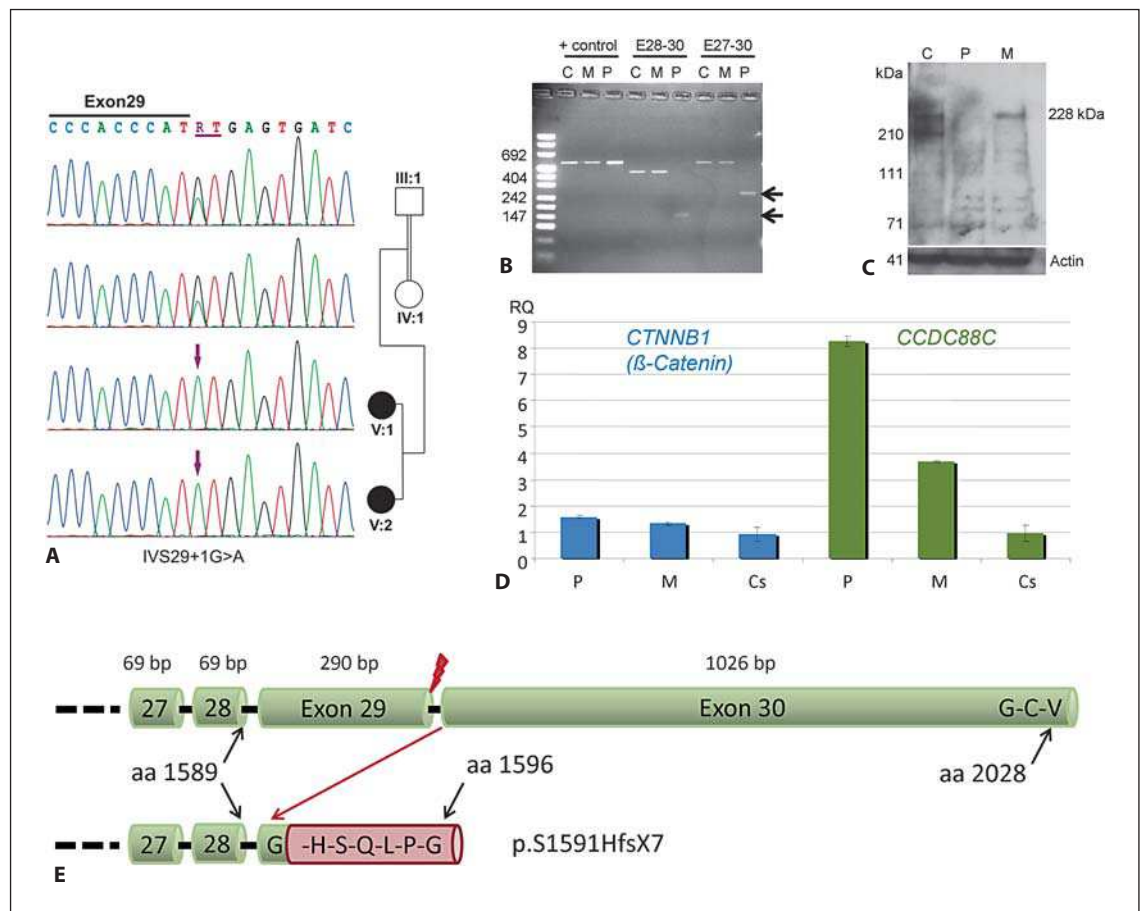


Fig. 4. Characterization of the detected splice site mutation IVS29 + 1G>A. C = Control, M = Mother, P = Patient. **A** Electropherograms of the homozygous mutation detected in both affected siblings and heterozygous carrier status of both parents. **B** Confirmation of an aberrant transcript by reverse transcription PCR. The aberrant splicing in the patient caused by the mutation in intron 29 results in deletion of exon 29, which was confirmed by sequencing of the aberrant transcript. Deletion of exon 29 (290 bp) results in a frameshift in the last exon and subsequent loss of the terminal Dishevelled binding site. **C** Western blotting using an antibody against the epitope encoded by exons 28 and 29 confirmed absence of wild-type protein in the patient. **D** Results of quantitative RT-PCR using minor groove binding probes for *CTNNB1* (β-catenin) and *CCDC88C* on cDNA derived from PAXgene RNA tube samples from the patient and her mother in

comparison to 8 unrelated healthy controls. In line with the assumed loss of function without nonsense-mediated mRNA decay, we found significantly increased expression of both *CCDC88C* and β-catenin in the patient. The patient's heterozygous mother showed also increased *CCDC88C* expression levels but no statistical significant alteration of β-catenin expression ($p < 0.028$ in the patient, $p < 0.148$ in the mother; Wilcoxon two-sample test). **E** Scheme of the effect of the splice site mutation on the protein level. Upper cartoon showing the terminal part of the wild-type protein ending with the G-C-V Dishevelled binding site. Lower cartoon indicates the frameshift resulting from loss of exon 29 after the first amino acid from exon 30, the addition of 6 amino acids and the loss of the Dishevelled binding site (p.S1591HfsX7). Red flash indicates site of splice site mutation.

transcription due to loss of *CCDC88C* function. β-catenin expression showed a significant increased expression in the patient only, while the expression level in the mother was not significantly altered (fig. 4D). Expression profiling for the Wnt pathway genes revealed significant down-regulation of many genes in the patient, some of which, though, were also diminished in the healthy heterozygous

mother (fig. 5). Genes that were significantly down-regulated in the patient, but not in the mother, were: *AXIN1*, *CSNK1D*, *CTBP2*, *EP300*, *FBXW2*, *FRAT1*, *FZD1*, *GSK3A*, *KREMEN1*, *SLC9A3R1* and *TCF7L1*. *LEF1* was the only gene that apart from β-catenin was significantly up-regulated in the patient and did not show a significant change in the mother (fig. 5, 6).

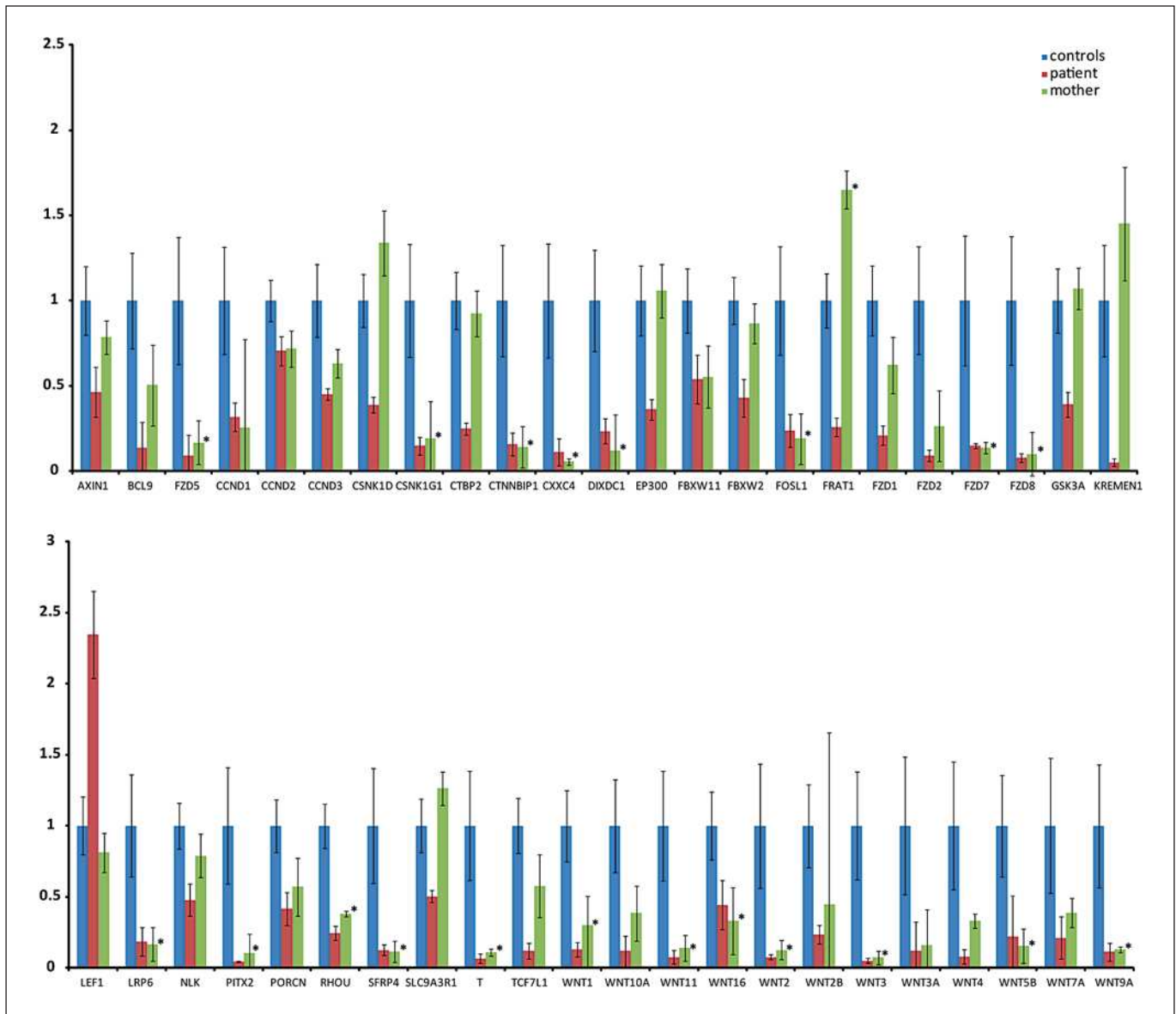


Fig. 5. Diagram of relative changes in statistically significant differentially expressed genes in blood of the patient (red) using the Human Wnt Signalling Pathway PCR Arrays (SABioscience). Values of the healthy carrier mother (green) that also show statistically significant alterations are marked with an asterisk. Note significant over-expression of LEF1 and down-regulation of many WNT signalling proteins including the Frizzled receptor and transcriptional target genes in the patient. Standard deviations in

the patient and the mother are derived from 4 different experiments each, while the standard deviations in controls are derived from single experiments in six different individuals. The commonly narrow standard deviations in the patient and her mother reflect the high reliability of the measurements, while the quite big standard deviations in the controls indicate a broad biological variability in expression levels for many of the genes in blood.

Expression Profiling of CCDC88C

In order to gain further insight into the developmental role of *CCDC88C* and its different isoforms in humans, we performed isoform-specific expression profiling by quantitative RT-PCR from cDNA panels of various hu-

man fetal and adult tissues. These analyses revealed highest fetal expression in the brain, especially in the pons and cerebellum, while in adult brain tissues expression was highest in the cortex. The highest expression in adult tissues was found in the pancreas. Expression was also rela-

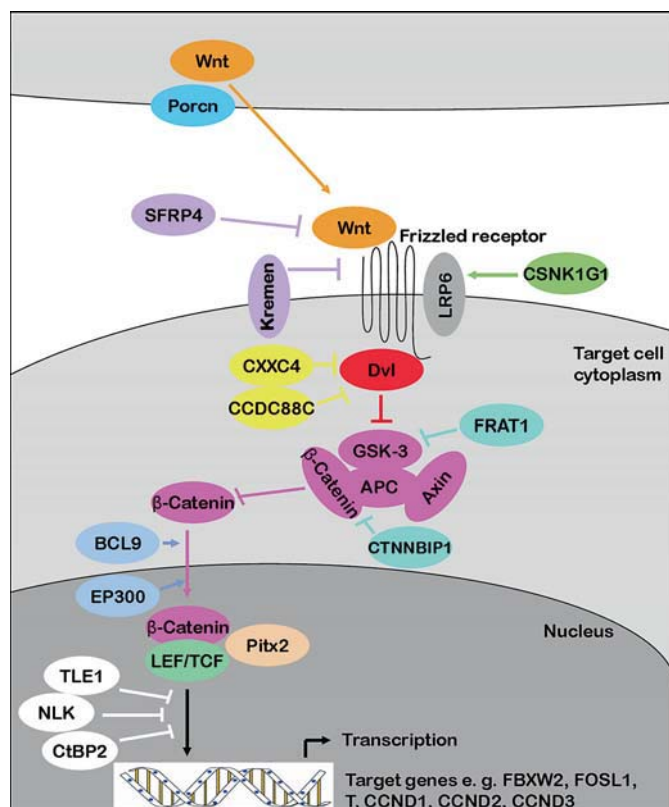


Fig. 6. Scheme of the Wnt signalling pathway including proteins from the RT-PCR array. Porcn, a transmembrane protein is involved in the secretion of Wnt proteins, which represent intercellular signalling molecules [Rocheleau et al., 1997]. Reaching their target cells, Wnt proteins bind to Frizzled transmembrane receptors. The secreted protein SFRP4 shows homology to the cysteine-rich ligand binding domain of the Frizzled receptors and thus functions as an antagonist of Wnt binding [Rattner et al., 1997]. Kremen also has an antagonistic function, since it represents the receptor of the negative regulator Dkk [Mao et al., 2002]. LRP6 represents a co-receptor of Wnt proteins, which is regulated by CSNK1G1 [Davidson et al., 2005]. Dvl, which is inhibited by CXXC4 and CCDC88C, mediates Wnt signalling via inhibition of GSK-3. GSK-3 is also inhibited by FRAT1 [Yost et al., 1998] and β -catenin is inhibited by CTNNBIP1 [Tago et al., 2000]. By inhibition of the β -catenin degrading GSK-3/APC/Axin complex β -catenin accumulates in the cytoplasm and is transferred to the nucleus, where it associates with LEF, TCF or Pitx2 proteins to activate transcription of target genes. BCL9 and EP300 represent transcriptional co-activators [Sun et al., 2000; Hoffmans et al., 2005] and TLE1, NLK and CtBP2 represent negative regulators [Chen et al., 1999; Furusawa et al., 1999; Ishitani et al., 2003].

tively high in fetal and adult kidney and in fetal thymus (fig. 7A). Isoform specific RT-PCR revealed the expected products for isoforms IF1 (exons 14–15), IF2 (exons 25b–29), IF3b (exons 26b–28), IF3f3ar (exons 26–27b) and IF4

(exons 2–3b) in blood (fig. 7B and data not shown). IF3a (exons 26b–27b) could not be amplified. Isoform-specific RT-PCR from human brain cDNA panels revealed that in adult brain only the longest isoform 1 was expressed and only in the cortex and medulla oblongata. In fetal brain isoform 1 was detected in the cortex cerebri, cerebellum, pons and spinal cord. In addition isoform 3b was expressed in fetal cerebellum and pons and in adult medulla oblongata (fig. 7B).

Discussion

Via genome-wide linkage analysis we revealed a homozygous *CCDC88C* splice site mutation as underlying a distinct autosomal recessive complex hydrocephalic brain malformation with impressive prenatal enlargement of the lateral ventricles and the posterior fossa, and a diverticulum-like pouch. Despite the severe prenatal cerebral phenotype the affected patient V:2 showed normal psychomotor development at the age of 3 years and 3 months after ventriculo-peritoneal shunting and besides one febrile seizure no further organic or neurologic abnormality. One aunt of this patient, who was also an offspring of a consanguineous mating, was reported to have been mildly mentally retarded. Assuming the same condition in her, this may indicate that V:2 might show mild intellectual disabilities only later in life, or that this aunt might have developed normally if her hydrocephalus had been detected and treated accordingly. The diverticulum-like pouch in V:2 was initially considered to resemble schizencephaly, which is defined as true clefts formed in the brain as the result of failure of development of the cerebral mantle in the zones of cleavage of the primary cerebral fissures, which must be lined by abnormal gray matter (microgyria) [Leventer et al., 2008]. Mutations in *EMX2*, which is also involved in Wnt signalling, was reported to cause schizencephaly in 1996, however, this association has not since been confirmed [Merello et al., 2008]. Nevertheless, we sequenced *CCDC88C* in 58 patients with a radiological diagnosis of schizencephaly with or without reported associated brain malformations, but could not detect any mutation, underscoring that the brain malformation observed in the presented consanguineous family is distinct from schizencephaly. We then sequenced 5 additional patients with variable unclassified brain malformations with enlarged lateral ventricles, but did not find disease causing mutations. Therefore, *CCDC88C* mutations seem to cause a distinct non-syndromic complex hydrocephalus with apparently

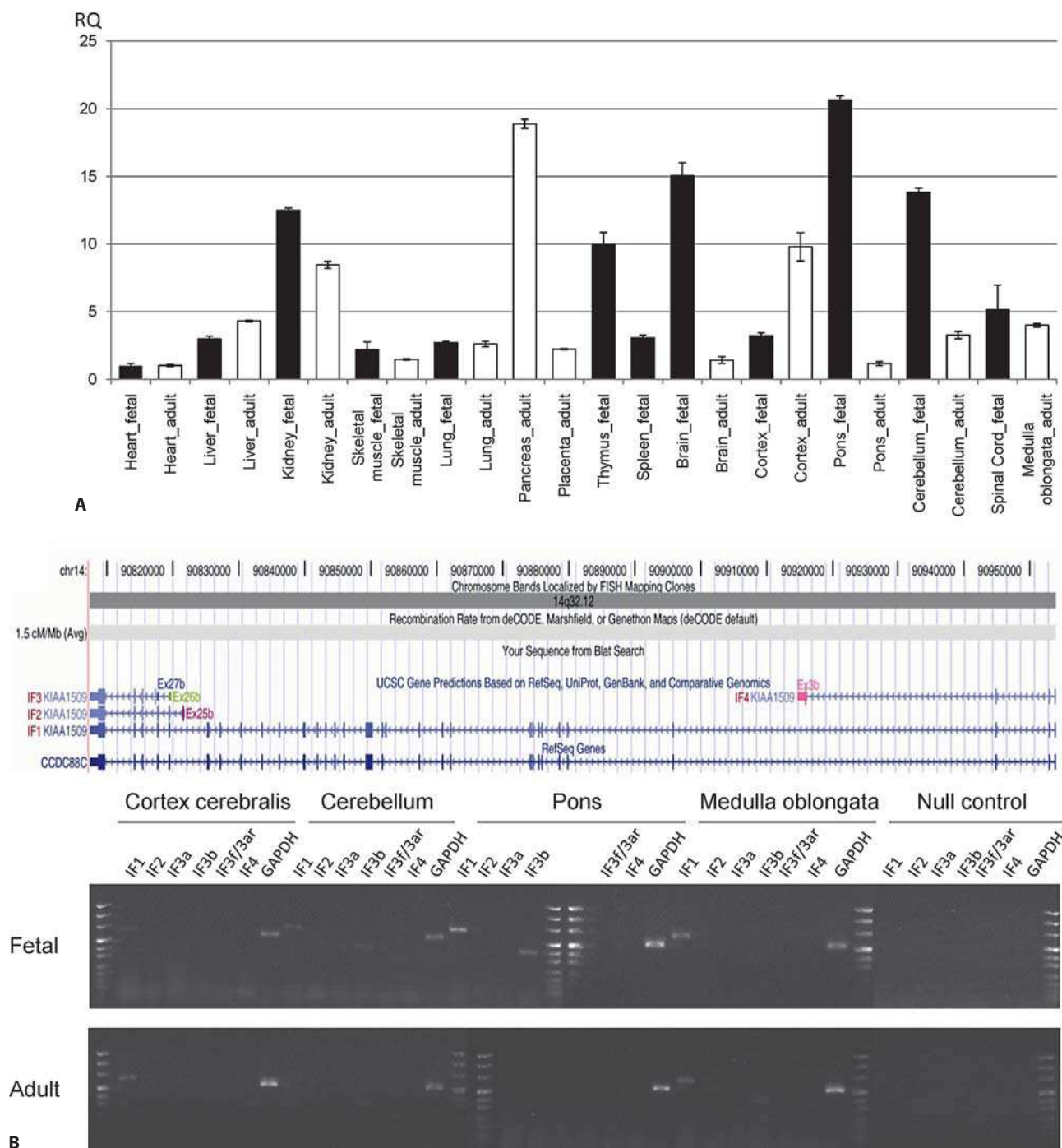


Fig. 7. *CCDC88C* expression studies by quantitative RT-PCR in cDNA from various human fetal and adult tissues showed highest relative expression in fetal brain and highest relative expression in adult pancreas (**A**). **B** Upper part represents a scheme of the various isoforms and the lower part shows the results of isoform-specific *CCDC88C* RT-PCR on fetal and adult brain cDNA panels. Size standard is puC Mix 8 (Fermentas) and GAPDH primers were used as positive control. IF1 is expressed in all fetal brain tis-

sues investigated, but only in adult cortex and medulla. From the other isoforms only IF3b was found expressed in brain tissues: in fetal cerebellum and pons and in adult medulla. Primers for isoform specific RT-PCR were located in exon 14/15 and exon 15 for IF1 (617 bp), exon 25b and exon 29 for IF2 (488 bp), in exon 26b and 27b for IF3a (403 bp), in exon 26 and 27b for IF3f/3ar (234 bp), in exon 26b and exon 28 for IF3b (540 bp) and in exon 2 and 3b for IF4 (540 bp).

normal psychomotor development despite the impressive prenatal cerebral phenotype.

CCDC88C encodes a member of the hook-related proteins also known as Hook-related protein 2 [Enomoto et al., 2006]. This protein family is characterized by a microtubuli binding aminoterminal domain and a central coiled-coil domain. Hook-related protein 2 has in addition a carboxy-terminal domain with residues Glycin-Cystein-Valin which can bind to the PDZ-domain of Dishevelled (Dvl), an important scaffold protein involved in the regulation of the Wnt signalling pathway. Therefore it is also known as DAPLE which stands for 'Dishevelled associating protein with a high frequency of leucin residues'. The Wnt protein family transmits a plethora of intercellular signals crucial for the development and homeostasis of metazoan animals [Logan and Nusse, 2004]. Dependent on the context, a given Wnt signal may denote cell proliferation, apoptosis, differentiation, migration, polarization, cell-fate specification, or other cellular processes. During early embryonal development WNTs are important caudalizing factors for the establishment of the initial anterior-posterior polarity of the brain. Disruption of Wnt-1 has been shown to disturb development of midbrain and anterior hindbrain structures including the cerebellum. Furthermore, an implication of Wnt signalling in the pathogenesis of hydrocephalus and cystic malformations of the posterior fossa is suggested by the observation of a complex phenotype including posterior cyst and postnatal hydrocephalus in transgenic mice with ectopically expressed *Engrailed-1*, a homeodomain transcription factor downstream of Wnt-1 [Rowitch et al., 1999]. Likewise, the distinct phenotype of the index patient in this study includes a hydrocephalus, a medial diverticulum and marked retrocerebellar fluid accumulation in an enlarged posterior fossa (fig. 1C).

Interestingly, cerebral midline cysts with or without enlarged ventricles are commonly observed as 'posterior cysts' within the phenotypic spectrum of holoprosencephaly and/or as 'interhemispheric cysts' in patients with agenesis of the corpus callosum [Simon et al., 2001], which assumingly may result from heterozygous mutations in the 2 genes *SIX3* and *SHH*. Both genes as well as *LICAM* have been shown to interact with Wnt signalling during early embryonal development [Lacbawan et al., 2009; Grana et al., 2010; Sylvester et al., 2010].

Wnt signals are not only crucial for development, but are also required for adult tissue maintenance including synaptic function, and perturbations promote human degenerative diseases and cancer. Wnt signalling is mediated through the seven-transmembrane Wnt receptor,

Frizzled, which transmits the signal downstream via the cytoplasmic protein Dvl. There are two distinct pathways downstream of Dvl, the canonical β -catenin pathway and the non-canonical JNK pathway. Binding of Wnt to Frizzled leads to the accumulation, nuclear translocation, and interaction of β -catenin with the Tcf/Lef family of transcription factors. Therefore, β -catenin functions as a co-activator of transcription of Wnt-responsive genes. In the mouse, *Daple* inhibits Wnt3a induced accumulation of β -catenin and activation of T-cell factor transcriptional activity [Oshita et al., 2003]. In contrast to murine *Daple*, the *Xenopus* homologue of *Daple* activates Wnt signalling during development [Kobayashi et al., 2005].

The splice site mutation in the presented family leads to a premature stop codon within the last exon of *CCDC88C*, therefore assumingly not resulting in nonsense mediated mRNA decay. This was confirmed by quantitative RT-PCR in the patient's blood sample which indeed showed significantly increased *CCDC88C* mRNA levels, probably as a feedback mechanism from loss of function. As expected from the loss of function of the negative regulator *CCDC88C* through failure to inhibit Dvl which itself inhibits the GSK3-APC complex which normally inactivates β -catenin, we also found increased levels of β -catenin and the downstream protein LEF1 in the patient. Nevertheless, many upstream and downstream inhibiting as well as promoting Wnt proteins showed reduced expression, which could be explained by a complex negative feedback loop (fig. 5, 6). Therefore, loss of human DAPLE might reduce Wnt signalling mediated gene transcription due to a negative feedback loop. However, since the healthy mother also showed diminished expression of some of the genes, expression studies in patient's blood may not reflect the mechanisms during development and neuronal maintenance. Therefore, further studies in model systems are necessary to elucidate these findings. Nevertheless, these data indicate that disturbed Wnt signalling would have been detectable as an underlying disease cause in the patient by initial expression profiling. Since expression profiling is increasingly considered as a screening method to narrow down the underlying cause in patients with clinically and genetically heterogeneous conditions, this expression profile may help to identify further patients with *CCDC88C* mutations.

The expression pattern of DAPLE in embryonic mammals has not been examined, but expression of *Xenopus Daple* was found in various developing tissues including the mesencephalic region of the neural plate, the tail, eye and otic vesicles, olfactory placode, and the pharyngeal

cavity [Kobayashi et al., 2005]. We investigated expression of human *CCDC88C* in a panel of embryonic and adult tissues by quantitative RT-PCR and found highest expression in fetal pons, cerebellum, thymus and kidney, as well as in adult pancreas, kidney and cortex. Nevertheless, loss of functional DAPLE in humans seems to affect only brain development. This finding may be related to isoform specific functions supported by our finding of nearly exclusive expression of isoform 1 in various fetal and adult brain tissues. The high *CCDC88C* expression in fetal pons and cerebellum corresponds well to the apparently abnormal development of the posterior fossa observed in our patient.

In conclusion, we report the first human phenotype associated with a mutation of the *CCDC88C* gene which encodes the Wnt signalling inhibitor DAPLE. Despite a dramatic prenatal hydrocephalic brain malformation, the affected living patient develops within the normal range and appears to have no further health issues.

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